#### TENT COOPERATION TREAL Y

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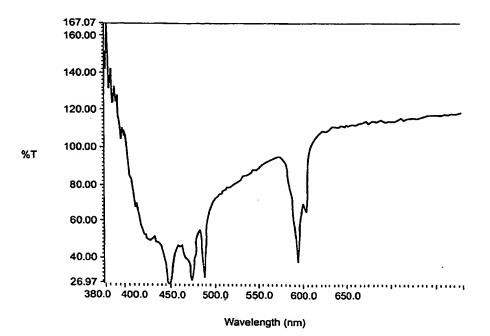
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#### (57) Abstract

A method of providing covert security features for documents such as vouchers, packaged goods and banknotes in which the document is provided with a dopant. The dopant consisting of a material which can be identified by examination of its response to visible wavelength photon radiation and which can be applied directly on or into the document or can be fused into glass matrices before application.

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SECURITY PRINTING 1 2 3 The invention relates to materials and techniques 4 relating to security printing. 5 6 The present invention in its broadest sense is 7 concerned with the provision of security in relation to 8 documents, vouchers, packaged goods and tokens of 9 value. Examples of these are banknotes, cheques and 10 drafts, bond and stock certificates, and credit and 11 bank cards. All of these are referred to hereinafter 12 for simplicity as "documents". 13 14 Documents of this nature have the requirement to be as 15 secure as possible against forgery and falsification 16 and for this purpose it is desirable that they exhibit 17 both covert and overt security features. 18 expression "covert security feature" is used to denote some security feature which is not visually apparent to 19 the normal user, whereas "overt security feature" is 20 21 used to denote a feature which can be readily seen and 22 recognised by members of the public without the use of

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1 specialised equipment or confidential information. 2 Traditional forms of overt security features include 3 water marks, metal security threads, and the use of 4 specialised forms of paper and printing. 5 6 Known methods of covert security include NIR and IR 7 absorber inks, magnetic threads, complex optical and electrically conductive indicia, anti-Stokes, visible-8 wavelength-emitting phosphors etc. 9 10 11 With rapid advances in reprographic technology such as 12 relatively cheap and high quality colour photocopiers and easily available digital image manipulation, the 13 14 traditional forms of security have become increasingly easy to circumvent. 15 This is because the absorption and 16 emission in the visible, NIR and IR ranges of all the 17 currently used and proposed security dopants are 18 readily available in the public domain since the current materials were developed for the laser and lamp 19 industries. This is particularly true for all the rare 20 21 earth containing absorbers and emitters, where many 22 thousands of public domain references of absorption and emission spectra are listed from the 1950's onwards. 23 24 There is accordingly a requirement for improved forms 25 of both covert and overt security features, preferably 26 ones which can be used with existing printing 27 technology at modest cost. 28 29 According to one aspect of the present invention, there is provided a method of providing a document with a 30

covert security feature, in which the document is

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printed using an ink containing a dopant, the dopant 2 being of a material which can be identified by 3 examination of its response to visible wavelength 4 photon radiation. 6 This and other aspects and features of the present 7 invention are defined in the appended claims. 8 9 The present invention will now be described by way of 10 example with reference to the accompanying drawings of 11 which: 12 13 Fig. 1 shows a blue ink reflectance spectrum from a 14 paper print; 15 16 Fig. 2 shows green ink reflectance spectrum from a paper 17 print; 18 19 Fig.3 shows red ink reflectance spectrum from a paper 20 print; 21 22 Fig.4 shows a reflectance spectrum from the 23 Praesodymium Oxide dopant in accordance with the 24 present invention; 25 26 Fig. 5 shows a reflectance spectrum from the Neodymium 27 Oxide dopant in accordance with the present invention; 28 29 Fig. 6 shows a reflectance spectrum from the Holmium 30 Oxide dopant in accordance with the present invention;

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1 2 Fig. 7 shows a reflectance spectrum from the Thulium 3 Oxide dopant in accordance with the present invention; 4 5 Fig.8 shows a reflectance spectrum of raw Europium 6 Oxide powder as used in the present invention; 7 8 Fig.9 shows a reflectance spectrum of the same 9 Europium Oxide contained in glass; 10 Fig.10 shows a reflectance spectrum of raw Erbium Oxide 11 12 powder as used in the present invention; 13 Fig.11 shows a reflectance spectrum of the same Erbium 14 15 Oxide contained in glass; 16 17 The present invention provides a range of inorganic 18 dopants designed with absorption spectra sufficiently 19 different in form and structure from the absorption 20 spectra of printing inks so that the dopants can be 21 easily identified. They thus become very covert because 22 they exhibit no UV, visible or IR stimulated output to 23 be observed by a counterfeiter. 24 25 The preferred elements for our dopants can be fused 26 with other elements in order to hide the presence of 27 the dopant element, or to alter its absorption 28 spectrum; or the oxide or salt of preferred element 29 itself can be directly mixed into, for example, a 30 printing ink or a batch composition for plastics production etc. When the dopant is mixed with other 31

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1 elemental compounds and where one of its admixture 2 compounds contains a substantial proportion by weight 3 of a particular range of atomic number (z) elements, 4 varying the proportion of this compound in the final 5 mix can vary the absorption spectrum of the final inorganic mixture, thus essentially creating further 6 7 dopants. 8 The present invention depends on the incorporation of a 9 synthesised inorganic dopant into or onto the document 10 at any stage of its manufacture, including the printing 11 12 These dopants are designed to have very complex 13 visible wavelength absorption spectra, measured in 14 either reflective or transmissive mode. The spectra 15 they exhibit are <u>not</u> found in printing inks or common marbling substrates. This results in high signal-to-16 17 noise ratio detection, and hence the ability to 18 identify the dopant in 10msec or less using low output 19 (c. 4W) bulbs as illuminants. 20 The dopant incorporation with its unique spectrographic 21 22 pattern gives independence from document soiling, wear 23 and tear etc, because it allows excellent signal-to-24 noise ratio. Pattern recognition software to identify, within 1 msec, the complex signature of our synthesised 25 26 dopants is readily available from suppliers in the 27 public domain, having been used in optical and nuclear 28 spectrometry for 30 years. Dopants in accordance with the present invention can be incorporated singly, 29 30 mixed, or in separate areas to produce a "bar code", or

to simply confuse a forger. The dopants, depending on

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1 composition, are either colourless or transparent, or 2 coloured, at the choice of the user. Dopants made in 3 accordance with the present invention provide high optical absorption yet give optical transparency 4 5 because their absorption features are created at 6 wavelengths to which the human eye is insensitive. 7 8 For visible wavelength interpretation the preferred 9 method is to illuminate an area of at least 5mm<sup>2</sup> by a 10 ring of at least 6-8 200µ optical fibres in a 11 concentric ring, and channel reflected light through an 12 inner 200µ optical fibre to the wavelength detector. It 13 has been found that this number of optical fibres gives 14 sufficient signal for interpretation of the spectra, 15 however the present invention is not limited to this 16 method of detection of the spectrum or the number or 17 arrangement of optical fibres used in this detection 18 method. This eliminates the optical losses due to 19 lenses in much prior art, which in turn leads to the 20 processing speed of our system. CCD based wavelength 21 detectors, followed by A-D conversion for processing 22 are standard technologies in public domain electronics. Our dopants are engineered to give no visible signal, 23 24 such as fluorescence, upon illumination by UV, visible, 25 or IR radiation and are hence not easily replicated as 26 has happened with fluorescent inks, and other emitting 27 technologies. 28 29 The advantages of the present invention will be readily

apparent when the spectra obtained from these dopants

7

is compared with those obtained from standard printing 1 2 inks, or colourisers in plastics etc. The standard inks and the like give relatively unsophisticated 3 reflectance spectra - see for example Figures 1, 2, 3. 4 These show the visible reflectance spectrum of a 5 Pantone standard blue, green and red ink from a paper 6 7 print. Figures 4, 5, 6, 7 show the visible reflectance spectra from the four dopants, Praesodymium Oxide, the 8 Neodymium Oxide, the Holmium Oxide and Thulium Oxide, 9 incorporated in a clear litho varnish and printed on 10 11 the same paper as that used to obtain the spectra shown in Figs. 1, 2 and 3. 12 13 The prints obtained using dopants in accordance with 14 15 the present invention are completely colourless to the 16 Figure 4 for example, shows many easily 17 identifiable peaks, troughs and turning points in its 18 spectrum with a shape easily distinguished from any ink or colouring dopants. It is these unique features 19 which give the excellent signal-to-noise ratio, giving 20 the rapid identification ability of our system, with 21 22 excellent identification rates, and very low false 23 acceptances, together with high rejection for forged 24 copies.

25

26 The features, and/or slopes, of the reflectance spectra can be shifted to create other dopants by incorporating 27 28 the dopants into inorganic compounds of the type 29 described later.

8

The use of visible wavelength spectrometry, as opposed 1 2 to IR or NIR wavelengths, makes possible many more 3 commercial applications. This is firstly because of the reduced cost of components for the visible, and 4 5 secondly because the cheapest excitation source is a 6 common (4W) torch bulb which emits plenty of visible 7 light but very little IR. Hence IR and NIR techniques 8 require more powerful and costly excitation sources. 9 Also by moving to the visible we make it easy to 10 construct simple hand-held portable instrumentation 11 which again increases possible commercial applications. 12 13 Visible wavelength spectroscopy as revealed in the 14 prior art with application to security uses lenses or 15 mirrors and lamps to provide the illumination source. 16 17 Many suppliers, such as Oriel Corp. USA, now make 18 commercially available reflectance probes which are about 6mm diameter overall and contain a ring of 19 20 illuminating fibres (200µ diameter 6-8 in number) 21 surrounding a centre core of detecting fibres. Use of 22 these probes gives much improved signal-to-noise ratio 23 at the CCD array, or Si photodiode array, or other 24 detector. Using other off-the-shelf components the 25 output of the array spectrometer can be coupled to D-A 26 converters and operated from a laptop, hand-held 27 palmtop, or desktop PC computers. This can easily be 28 interfaced to standard computer software on production 29 lines for authentication at high speed - 10m/sec.

9

1 The dopants we have identified as working well can be 2 added to standard offset litho printing inks in a manner known to those skilled in the art. 3 4 in quantities up to about 30% by volume without 5 affecting the printing process, providing the dopants 6 have been micronised into fine powders of the order of 7  $1-4\mu m$  diameter. If this step is omitted poor uniformity printing results. Our dopants need add no 8 9 colour to the ink, so give a colourless invisible 10 printed strip onto the object to be protected. 11 Alternatively a colouring dopant can be selected to 12 blend in with an existing coloured scheme. 13 14 A major advantage of the dopants made in accordance 15 with the present invention is that they are cheap and 16 simple, not requiring the presence of complex expensive 17 chemicals. 18 19 The dopants can be applied to artefacts by any standard 20 deposition technique - air spray, lacquering, printing, 21 stamping. 22 23 The dopants could also be directly incorporated into paper or plastic (for example) at time of manufacture 24 25 of said paper or plastic. For our techniques to work 26 it is not necessary that the dopants are added as a

30 require surface deposition can offer more

will be the simplest and cheapest method.

27

28

29

31 security/covertness to the process. It arises because

superior layer or film, although in many cases this

that our dopant/excitation/detector technology does not

10

1 the excitation methods we are employing have ranges of many tens of microns in common materials such as paper 2 3 and plastics. Since dopants in accordance with the 4 present invention need not be on the surface of the 5 document the forger is denied the opportunity to scrape off samples from repeated small surface areas and 6 7 analyse them to look for "surprising" changes in composition from area to area. Such changes give the 8 9 forger a clue that covert technology is being used in 10 that area. The multiple peaks, troughs, and turning points 11 12 resulting give rapid, positive, unambiguous 13 identification of dopant presence (and hence object 14 authenticity) and allow multiple dopants to be used as 15 a further method of disguise, if required. 16 17 The preparation of the inorganic powders for doping to 18 permit identification by visible light is not limited to the use of chemical compounds which could be formed 19 20 by precipitation from a solution because such compounds are limited in numbers. 21 It has been found that the 22 most useful compounds (those with the most distinctive 23 absorption spectra in the visible) could be formed by fusion melting. Silicates, phosphates, borates have 24 25 been found to be the most useful starting points for 26 fusion, because they give transparent glass matrices. 27 In forming the required solids for powdering, the

28

29 chemical batch composition is not, for example, limited

30 to that required to produce, say, a glass.

because long range atomic order is not required in the 31

11

1 solid, since homogeneity is assured by micronising the 2 composition. Indeed in general terms we have found 3 that the best compositions are obtained where phase 4 separation of the melt temperature is imminent. 5 point is determined experimentally for each 6 composition. Nor need the chemistry be limited to 7 stoichometric ratios such as to arrive at crystalline 8 compounds, e.g. as used to produce the commonplace 9 inorganic fluorescence powders added to printing inks. 10 11 In many compositions, the structure and magnitude of the absorption peaks can be controlled over a wide 12 13 range by control of the gas atmosphere during the melt This is established by trial and error for each 14 15 composition by test melting each composition in air, in 16 a reducing atmosphere, and in an oxidising atmosphere 17 to determine the optimum methodology and conditions for 18 the absorption profile required. 19 20 In many compositions, the structure and magnitude of 21 absorption peaks can be controlled by including a 22 substantial quantity (>20% by weight) of a high atomic 23 number Z element in the batch composition (lanthanum, 24 bismuth, and strontium work well, as examples). 25 varying the content of this high Z element only gives 26 changes in position and magnitude of the absorption 27 peaks, from composition to composition. Different 28 absorption peak wavelengths and magnitudes from that 29 exhibited by the raw dopant before being incorporated in a glass. 30

12

1 The effect of incorporating the dopant in a glass on

2 its spectrum can be seen in Figs. 8, 9, 10 and 11.

3

4 Fig. 8 shows a plot of the percent transmission against

- 5 wavelength (nm) for a raw Europium Oxide dopant powder.
- 6 Fig.9 shows a plot of the percent transmission against
- 7 wavelength (nm) for a Europium Oxide dopant powder
- 8 incorporated in a glass and ground into a fine powder.
- 9 The substances contained in the glass are as given in
- 10 Table 1 below and the glass plus dopant is made in
- 11 accordance with the method given below Table 1 on page
- 12 14.

13

- 14 Simply from a visual inspection it can be seen that the
- 15 two spectra are very different.
- 16 The feature of the spectrum of Europium Oxide shown at
- 17 reference numeral 81 for the raw oxide powder that
- 18 occurs at a wavelength of 533 nm has been shifted to
- 19 531nm. A similar shift can be seen for the lower
- 20 wavelength peaks 83 and 93. In both cases, the shift
- 21 in wavelength was 2nm. The most significant difference
- 22 between the spectra of Fig. 8 and Fig. 9 is the presence
- 23 of the line in the spectrum of the Europium Oxide
- 24 contained in glass at 393nm. There is no similar line
- 25 in the raw powder spectrum.

- 27 Fig. 10 shows a plot of the percent transmission
- 28 against wavelength (nm) for a raw Erbium Oxide dopant
- 29 powder. Fig.11 shows a plot of the percent
- 30 transmission against wavelength (nm) for an Erbium
- 31 Oxide dopant powder incorporated in a ground fine

13

1 powder glass. As with the sample used to obtain the 2 spectrum if Fig.9, the substances contained in the 3 glass are as given in Table 1 below and the glass plus 4 dopant is made in accordance with the method given 5 below Table 1 on page 14. 6 7 Fig. 10 shows, at reference numeral 101, the existence 8 of multiple peak structure occurring from a minimum 9 point at 654nm to approximately 700nm. It can be seen that these features are absent from the spectrum of 10 11 Fig. 11 as indicated at reference numeral 111. 12 13 Fig. 10 also has multiple peak structure occurring from 14 a minimum value at 521nm up to approximately 600nm. 15 These features are absent from the spectrum of Fig. 11 16 as can be seen at reference numeral 113. 17 18 We have shown our dopant technology to work in a wide 19 variety of compounds, including, but not limited to, 20 silicates, borosilicates, borates and germanates. 21 22 The following are a number of examples of the 23 composition and method of manufacture of a doped glass 24 in accordance with the present invention. 25 26 Example 1 27 28 A glass batch of a typical suitable composition is as 29 follows.

14

1

2

Compound	Wt %
SiO <sub>2</sub>	35%
B <sub>2</sub> O <sub>3</sub>	40.0
Na <sub>2</sub> O	8.5
K <sub>2</sub> O	8.5
Al <sub>2</sub> O <sub>3</sub>	1.0
Mg0	4.0

3

4

Table 1

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To this batch was added 0.1 to 25 wt% of Eu<sub>2</sub>O<sub>3</sub>. All 7 powder sizes can be used but approximately 250 mesh is 8 preferable. A wide range of crucibles can be used, a 9 Platinum crucible was used in this case. The final 10 batch is mixed and homogenised then it is added to the crucible heated to 845°C. The temperature is then 11 12 increased at a rate of approximately 5 °C/min to 1200 °C 13 the final melt temperature. It has been found that 14 good quality melts are produced by holding the melt at 15 the final temperature for between 2 and 2.5 hours 16 before powdering the glass. For absorber products not 17 visible to the naked eye, the natural emissions of Eu<sub>2</sub>O<sub>3</sub> 18 may be quenched by the use of high concentrations of 19 Eu<sub>2</sub>O<sub>3</sub> or by the inclusion of small < 1% quantities of 20 nickel oxide, silver oxide or lead oxide as 21 luminescence quenchers.

## 1 The following compositions may also be used

Compound	Wt (g)	Compound	Wt (g)	Compound	Wt (g)
S i O₂	55	SiO <sub>2</sub>	70	SiO <sub>2</sub>	50
B <sub>2</sub> O <sub>3</sub>	65	B <sub>2</sub> O <sub>3</sub>	80	Be₂CO₃	20
Na <sub>2</sub> CO <sub>3</sub>	29	Na <sub>2</sub> CO <sub>3</sub>	29	SrCO <sub>3</sub>	20
K <sub>2</sub> CO <sub>3</sub>	20	K <sub>2</sub> CO <sub>3</sub>	20	Na <sub>2</sub> CO <sub>3</sub>	10
Li <sub>2</sub> CO <sub>3</sub>	5	Li <sub>2</sub> CO <sub>3</sub>	5	K <sub>2</sub> CO <sub>3</sub>	10
Al <sub>2</sub> O <sub>3</sub>	2	Al <sub>2</sub> O <sub>3</sub>	2	Li <sub>2</sub> CO <sub>3</sub>	5
MgO	8	MgO	5	Al <sub>2</sub> O <sub>3</sub>	2
				MgO	5

Table 2

Compound	Wt (g)	Compound	Wt (g)
SiO <sub>2</sub>	35	SiO <sub>2</sub>	55
$B_2O_3$	80	B <sub>2</sub> O <sub>3</sub>	65
		<u> </u>	
Be <sub>2</sub> CO <sub>3</sub>	40	Na <sub>2</sub> CO <sub>3</sub>	29
Na <sub>2</sub> CO <sub>3</sub>	29	K <sub>2</sub> CO <sub>3</sub>	20
		i	
K <sub>2</sub> CO <sub>3</sub>	20	Li <sub>2</sub> CO <sub>3</sub>	5
Li <sub>2</sub> CO <sub>3</sub>	5	Al <sub>2</sub> O <sub>3</sub>	2
Al <sub>2</sub> O <sub>3</sub>	2	MgO	8
		]	-
MqO	8		··

Table 3

16

1 Another suitable composition is of the type

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3

Compound	Wt %
SiO <sub>2</sub>	51
B <sub>2</sub> O <sub>3</sub>	13
Al <sub>2</sub> O <sub>3</sub>	8
MgO	6
Ca0	10
Sr0	4
ZnO	4

4

5 Table 4

6

- 7 This is particularly suitable as a base for
- 8 incorporating dopants for visible wavelength absorption
- 9 detection because all the base elements have largely
- 10 unfeatured absorption spectra.

11

- 12 Dopants have also been successfully incorporated into
- 13 glass matrices with the following ranges of chemical
- 14 composition.

15

- 16 30-56wt% SiO<sub>2</sub>,
- 17 5-35wt%,  $La_2O_3/Bi_2O_3/Sr_2O_3$ ,
- 18 2-33wt%  $Li_2O/K_2O/Na_2O$ ,
- 19 0-6% Al<sub>2</sub>O<sub>3</sub>
- 20 wherein the La, Bi, Sr are examples of a suitable high
- 21 Atomic number component.

17

1 Incorporation of all three alkaline earth compounds, plus BaO, gives much reduced melting temperatures. 2 3 Preferred elements for dopant fabrication for visible 4 5 wavelength absorption system 6 7 Barium Zinc 8 Lanthanum Samarium 9 Lead Praesodymium 10 Magnesium Europium 11 Strontium Boron-10 12 Titanium Neodymium 13 Chromium Holmium 14 Iron Thulium 15 Caesium Cadmium 16 Molybdemum Antimony 17 Nickel Erbium 18 Tungsten Lutecium Cobalt 19 Tin 20 Sodium 21 Potassium 22 Terbium 23 Table 5 24 25 Improvements and modifications may be incorporated 26 without deviating from the scope of the invention.

18

1 CLAIMS

2

3 1. A method of providing a document with a covert

- 4 security feature in which the document is provided with
- 5 at least one dopant, the dopant being of a material
- 6 which can be identified by examination of its response
- 7 to visible wavelength photon radiation.

8

- 9 2. A method of providing a document with a covert
- 10 security feature as claimed in Claim 1, in which the
- 11 dopant comprises one or more inorganic compounds.

12

- 13 3. A method of providing a document with a covert
- 14 security feature as claimed in Claim 1 or Claim 2, in
- which the dopant comprises one of, or a combination of
- 16 the elements listed in Table 5, in elemental form or as
- 17 an oxide or salt.

18

- 19 4. A method of providing a document with a covert
- 20 security feature as claimed in any preceding Claim, in
- 21 which the dopant is mixed with a quantity of an element
- or its salt or its oxide with an atomic number greater
- 23 than 36.

24

- 25 5. A method of providing a document with a covert
- 26 security feature as claimed in Claim 4 in which the
- 27 element or its salt or its oxide is Strontium,
- 28 Lanthanum or Bismuth.

29

19

1 A method of providing a document with a covert 2 security feature as claimed in any preceding Claim, in which the dopant is mixed with ink and the resulting 3 4 mixture is applied to the document. 5 6 7. A method of providing a document with a covert security feature as claimed in any preceding Claim in 7 8 which the dopant is fused in a glass before being 9 applied to the document. 10 A method of providing a document with a covert 11 12 security feature as claimed in Claim 7 in which the 13 glass is made of silicates and/or phosphates and/or 14 borates. 15 16 9. A method of providing a document with a covert 17 security feature as claimed in Claim 7 or Claim 8 in 18 which the or each dopant is micronised into a fine 19 powder. 20 21 A method of providing a document with a covert 22 security feature as claimed in one of Claim 7 to Claim 23 9 in which each particle of the micronised fine powder 24 has a diameter of  $1-4\mu m$ . 25 26 11. A method of providing a document with a covert 27 security feature as claimed in any preceding Claim in which the dopant is such that, when the document is 28

illuminated with broad-band visible light to produce a

reflectance spectrum with frequency components

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- 1 generated by the dopant and by other reflecting
- 2 substances contained in the document, said spectrum
- 3 containing minimal frequency overlap between the
- 4 components of the spectrum generated by the dopant and
- 5 that part of the spectrum generated by other substances
- 6 contained in the document.

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- 8 12. A method of providing a document with a covert
- 9 security feature as claimed in any preceding Claim in
- 10 which the dopant is such that, when the document is
- illuminated with broad-band visible the frequency
- 12 components generated by the dopant are invisible to the
- 13 human eye.

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- 15 13. A method of providing a document with a covert
- 16 security feature as claimed any preceding Claim in
- 17 which the spectrum of the dopant can be shifted to a
- 18 higher or lower wavelength.

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- 20 14. A method of providing a document with a covert
- 21 security feature as claimed in any preceding Claim in
- 22 which the spectrum of the dopant can be shifted to a
- 23 higher or lower wavelength by alteration of the
- 24 composition of the glass in which it is fused.

- 26 15. A method of providing a document with a covert
- 27 security feature as claimed in any preceding Claim, in
- which the spectrum of the dopant is alterable by
- 29 alteration of the reaction temperature and/or pressure
- 30 at which the glass is made.

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1 2 16. A document provided with a covert security feature 3 by the method any of the preceding Claims. 4 A dopant for use in providing a document with a 5 covert security feature, comprising one or more 6 combination of the elements listed in Table 5, in 7 8 elemental form or as an oxide or salt, in finely divided form. 9 10 A method of making a dopant, in which one or a 11 12 combination of the elements listed in table 5, in elemental form or as an oxide or salt, is fused in a 13

glass and subsequently micronised.

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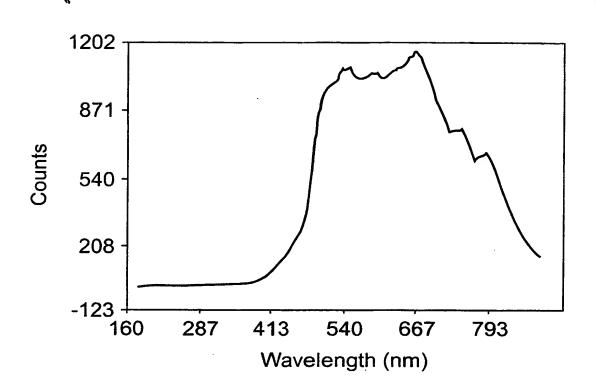


Fig. 1

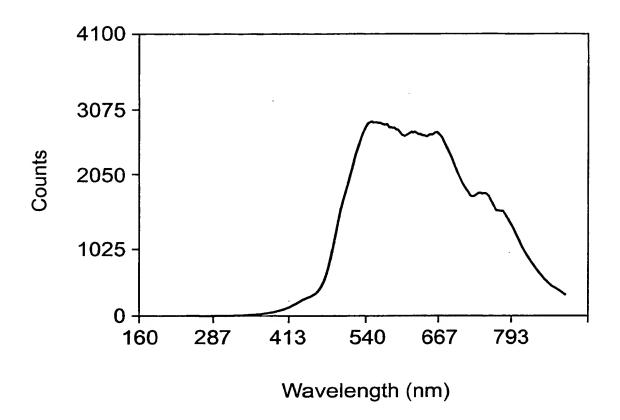


Fig. 2

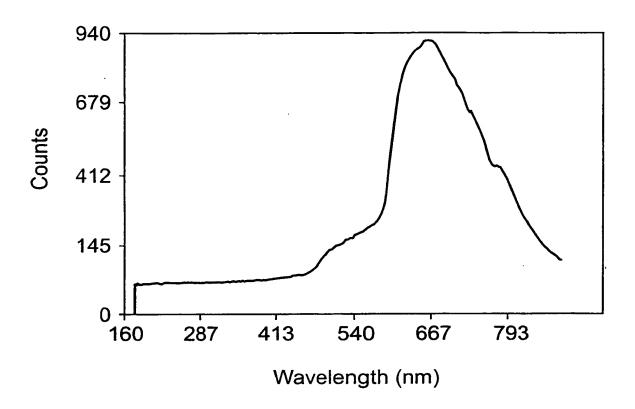
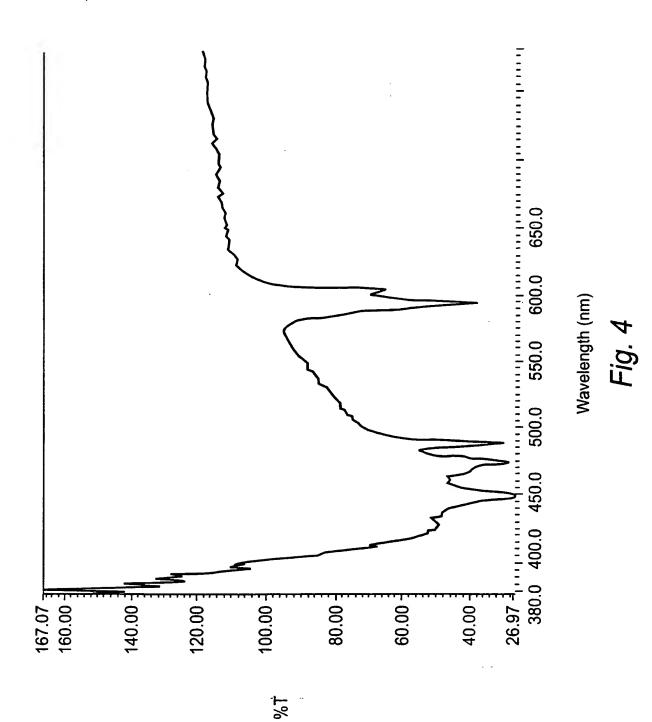


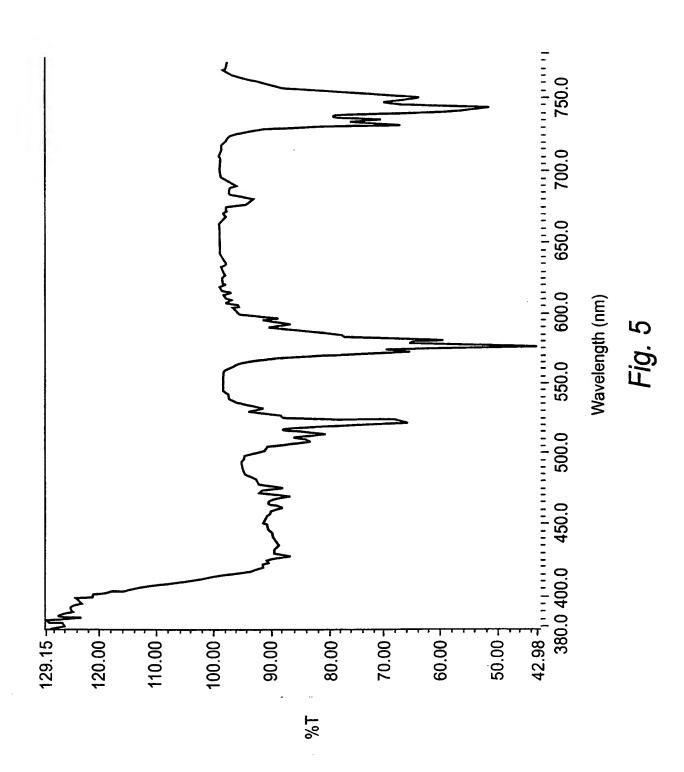
Fig. 3



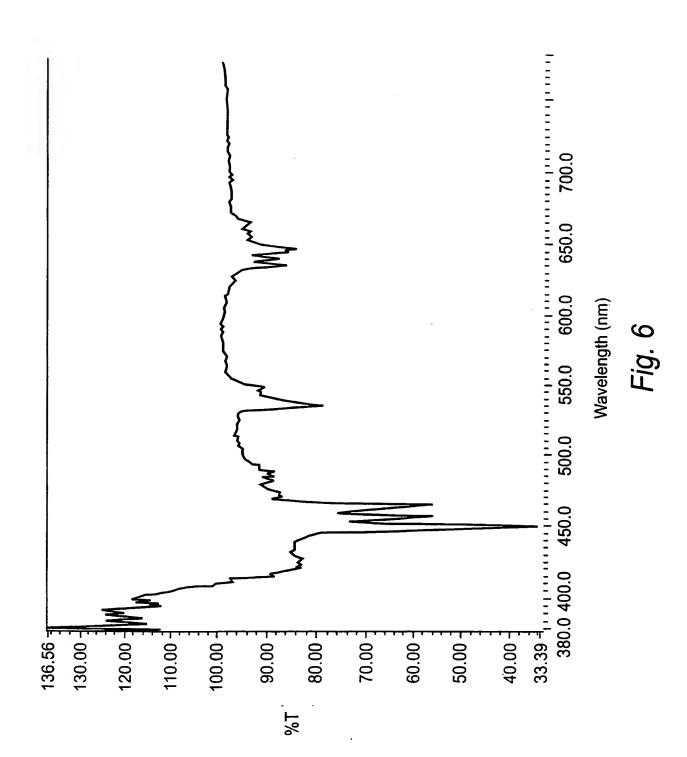
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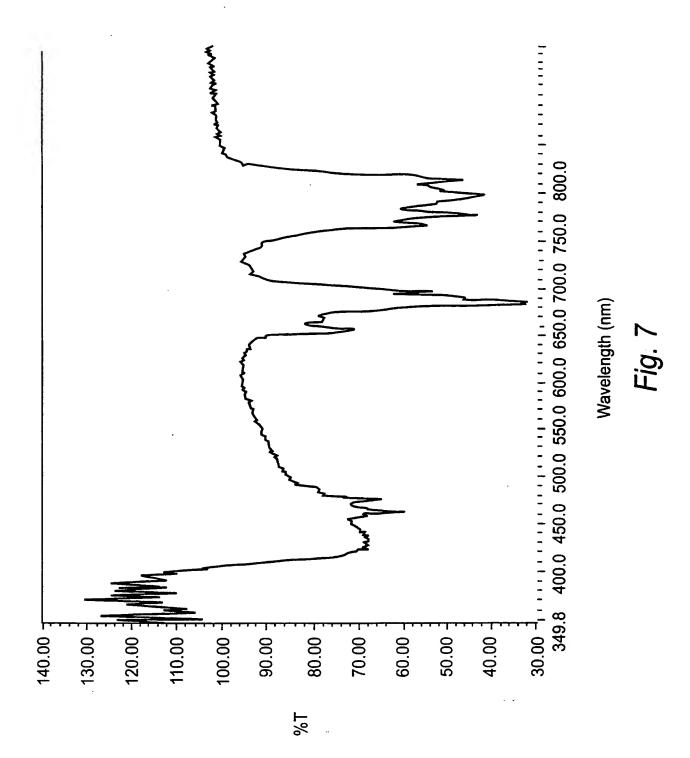


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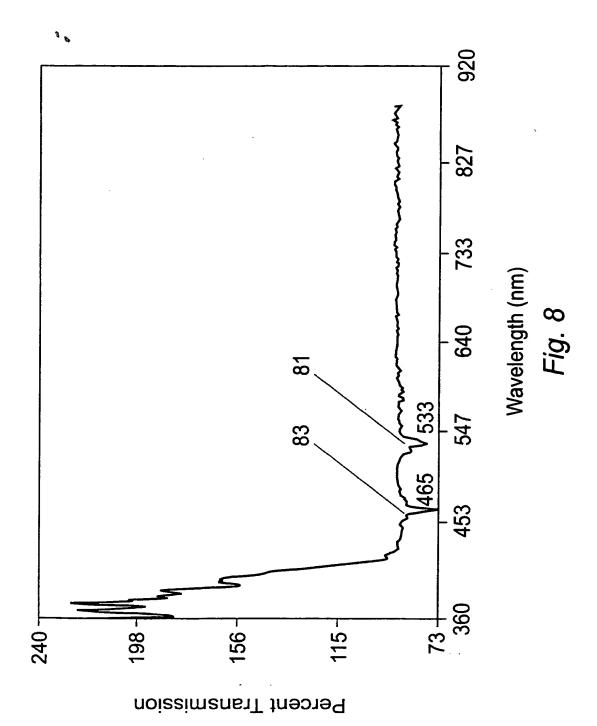
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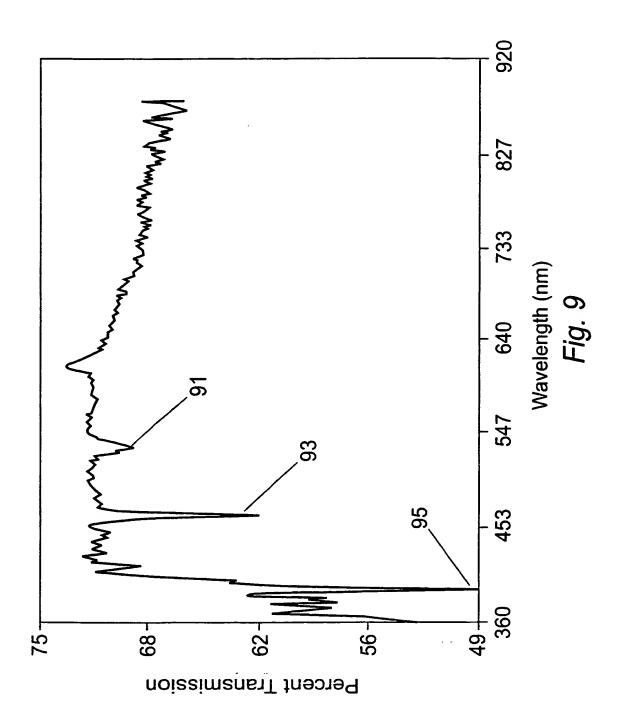


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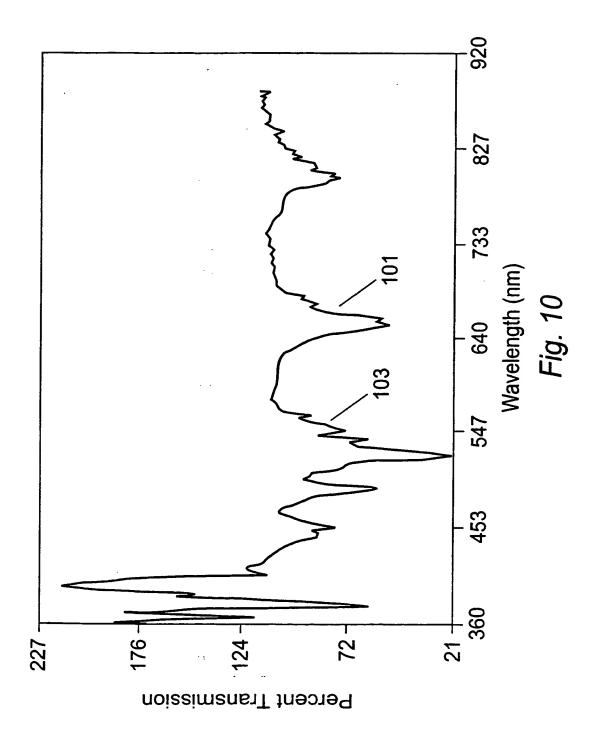
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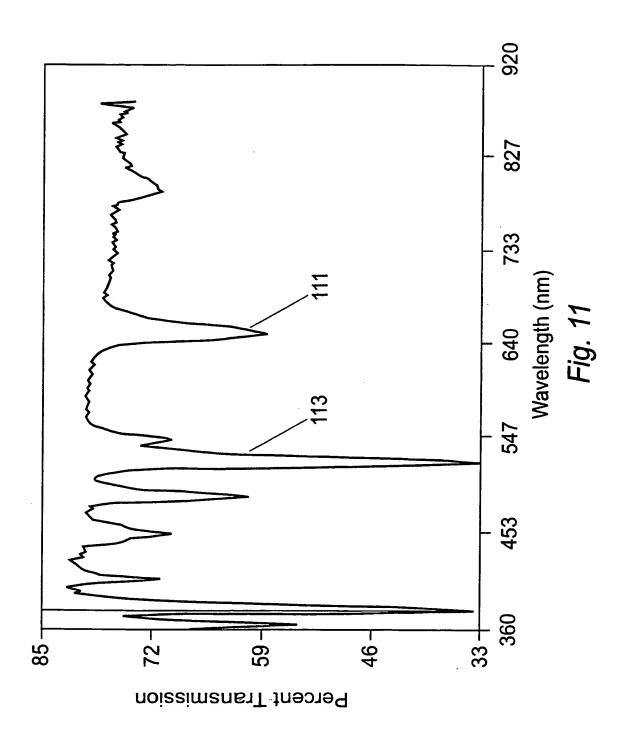
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Documenta	tion searched other than minimum documentation to the extent that s	such documents are inclu	ded in the fields se	arched
	lata base consulted during the international search (name of data ba	se and, where practical,	search terms used)	
	ENTS CONSIDERED TO BE RELEVANT			
Category °	Citation of document, with indication, where appropriate, of the rel	levant passages		Relevant to claim No.
X	WO 81 03509 A (GAO) 10 December 1981 (1981-12-10)  page 7, line 9 -page 8, line 5 page 11, line 17 - line 31 page 13, line 17 - line 29 page 14, line 21 -page 15, line 3 page 19, line 11 - line 13; clain	3 ns 9,10		1-5,7, 9-14, 16-18
X	EP 0 440 554 A (ARJOMARI EUROPE) 7 August 1991 (1991-08-07) page 5, line 60 -page 6, line 19;	; claim 6 -/		1-3,6, 11-13,16
X Furth	ner documents are listed in the continuation of box C.	χ Patent family π	nembers are listed in	n annex.
"A" docume conside "E" earlier of filing de which i citation "O" docume other n docume later th	and defining the general state of the art which is not bened to be of particular relevance to be of particular relevance to be of particular relevance to comment but published on or after the international state on the may throw doubts on priority claim(s) or so cited to establish the publication date of another or other special reason (as specified) on the referring to an oral disclosure, use, exhibition or neans of published prior to the international filing date but	cited to understand invention  "X" document of particul cannot be consider involve an inventive  "Y" document of particul cannot be consider document is combir ments, such combir in the art.  "&" document member of	not in conflict with the the principle or the dar relevance; the classification of the dar relevance; the classification of the dar relevance; the classification of the dark one or mornation being obvious of the same patent fa	ne application but boy underlying the simed invention be considered to urment is taken alone aimed invention entive step when the e other such docu- is to a person skilled
	3 January 2000	Date of mailing of the 08/02/20	ne international sear	ch report
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C (Continu	Intion) DOCUMENTS CONSIDERED TO BE COLOUR	FC1/GB 99/03092
Category :	Citation of document, with indication where appropriate, of the relevant passages	Relevant to claim No.
X Y	EP 0 211 754 A (PETREL) 25 February 1987 (1987-02-25) page 3, line 28 - line 34 page 4, line 21 - line 27 page 5, line 1 - line 24 page 6, line 13	1-3,6, 11-13,16 8,15
X	FR 2 568 191 A (OIFFER BOMSEL) 31 January 1986 (1986-01-31) page 5, line 15 - line 18; claims 1,2,8	1-3,6, 11-13,16
<b>X</b>	EP 0 202 902 A (ROYAL DOULTON) 26 November 1986 (1986-11-26) page 6, line 1 - line 8; claims 1,11,16	1-3,6-8, 16-18
X	DE 20 48 853 A (XEROX) 15 April 1971 (1971-04-15) page 5, line 21 -page 6, line 8	1-3
X	DE 28 45 401 B (GAO) 14 February 1980 (1980-02-14) claims 1,2,7,8,11,13	1-3
X	EP 0 037 507 A (AGFA-GEVAERT) 14 October 1981 (1981-10-14) page 5, line 21 -page 6, line 9; claims 1,4,5	1-3
X Y	EP 0 849 231 A (CORNING) 24 June 1998 (1998-06-24) column 1, line 14 - line 22; claim 1; figure 2	17,18 8,15
•		
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PCT/GB 99/03692

Patent document Publication Patent family **Publication** cited in search report date member(s) date WO 8103509 Α 10-12-1981 378210 B AT 10-07-1985 AT 900381 A 15-11-1984 CH 656655 A 15-07-1986 DE 3121523 A 15-04-1982 EP 0053148 A 09-06-1982 ES 503112 A 01-04-1982 FR 2484921 A 24-12-1981 GB 2088920 A,B 16-06-1982 IT 1144577 B 29-10-1986 JP 2012198 B 19-03-1990 JP 57500923 T 27-05-1982 450778 B SE 27-07-1987 SE 8107669 A 21-12-1981 US 4442170 A 10-04-1984 US 4533244 A 06-08-1985 EP 440554 07-08-1991 Α FR 2657630 A 02-08-1991 FR 2662717 A 06-12-1991 DE 69127481 D 09-10-1991 EP 211754 25-02-1987 FR 2585987 A 13-02-1987 **AT** 63500 T 15-06-1991 US 4736425 A 05-04-1988 FR 2568191 31-01-1986 NONE EP 202902 Α 26-11-1986 GB 2175851 A,B 10-12-1986 DE 2048853 Α 15-04-1971 GB 1332185 A 03-10-1973 DE 2845401 Β . 14-02-1980 CH 649788 A 14-06-1985 FR 2439094 A 16-05-1980 GB 2035208 A,B 18-06-1980 GB 2107646 A.B 05-05-1983 JP 55099000 A 28-07-1980 449069 B SE 06-04-1987 7908620 A SE 19-04-1980 SE 8401138 A 01-03-1984 US 4591707 A 27-05-1986 US 4691940 A 08-09-1987 EP 37507 Α 14-10-1981 DE 3013238 A 08-10-1981 AT 5957 T 15-02-1984 CA 1151225 A 02-08-1983 150881 A DK 04-10-1981 ES 501059 A 01-01-1982 JP 56151598 A 24-11-1981 NO 811141 A,B, 05-10-1981 72731 A,B PT 01-04-1981 US 4371196 A 01-02-1983 EP 849231 Α 24-06-1998 ΑU 4835597 A 25-06-1998 BR 9706433 A 25-05-1999 CA 2225138 A 20-06-1998 CN 1197217 A 28-10-1998 JP 10274718 A 13-10-1998